

Emergent properties hidden in plane view: Strong electronic correlations at oxide interfaces

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Complex oxides with correlated carriers are a class of materials characterized by multiple competing and nearly degenerate ground states due to interactions that create a subtle balance to define their ground state. This in turn leads to a wide diversity of intriguing properties ranging from high T_c superconductivity to exotic magnetism and orbital phenomena. By utilizing bulk properties of these materials as a starting point, interfaces between different classes of correlated oxides offer a unique opportunity to break the fundamental symmetries of the bulk and alter the local environment. From experimental point of view, utilizing recent advances in growth with atomic layer precision one can now combine layers of compounds with distinct and even antagonistic order parameters to design new artificial quantum materials. Here we illustrate this approach by selected examples how broken lattice symmetry, strain, and altered chemical and electronic environments at the correlated interfaces can provide a unique platform to manipulate this subtle balance and enable quantum many-body states and phenomena not attainable in the bulk.

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I. INTRODUCTION

Finding new collective electronic states in materials is one of the fundamental goals of condensed matter physics. While the traditional approach has been to search for such phases within naturally occurring compounds, in recent years the focus has shifted to *heterostructures* (Hwang

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et al., 2012): artificial materials formed by interleaving two or more structurally and chemically dissimilar materials. Of particular interest is the spatial region at the *interface* where dissimilar materials meet. New states may emerge here because the environment near an interface is different from that occurring in bulk (thermodynamically stable) materials. Advances in the *angstrom-scale* layer-by-layer synthesis of multi-element compounds for *materials-by-design* have taken the approach to a new level of power and sophistication: It enables the atomic-scale combination of materials with different properties, granting access to a new terrain in which unusual states of matter may arise (Schlom *et al.*, 2008).

Heterostructures formed from transition metal oxides (TMO) are a particularly appealing hunting ground for new physics. In these materials the transition metal (M) ion has an open d -shell electronic configuration with spin, orbital, and charge degrees of freedom. Electrons in these partially filled d -shells are *correlated*: the motion of one electron depends explicitly and non-trivially on the behavior of all of the others giving rise to interesting many-body phenomena (Imada *et al.*, 1998). The resulting magnetic, superconducting, and multiferroic phases are of great scientific interest and are potentially capable of providing innovative energy, security, electronics and medical technology platforms. The heterostructure geometry (Granozio *et al.*, 2013; Mannhart and Schlom, 2010; Zubko *et al.*, 2011a) enables otherwise unattainable changes in atomic structure and chemical bonding, leading to new modalities for control and optimization of known states and potentially leading to new ones.

Over the past decade, one particular class of heterostructures, based on the interface between lanthanum aluminate (LaAlO_3 ; LAO for short) and Strontium titanate (SrTiO_3 ; STO for short), has been the subject of very extensive study. In this Colloquium we choose not to discuss the LAO/STO interface or its variants, selecting our examples instead from vanadate, manganite, cuprate and nickelate-based systems for two reasons. First, the LAO/STO system and its variants have been extensively reviewed in other venues, see for example Hwang *et al.*, 2012; Mannhart *et al.*, 2008; and Zubko *et al.*, 2011b. Second, and more importantly, the LAO/STO system involves doping nominally insulating STO with maximum sheet carrier densities of fewer than 0.5 electrons (e) per in-plane unit cell, and the charge density is typically spread over several unit cells in the direction perpendicular to the interface. The volume carrier densities are therefore typically low, so that the situation is more closely related to a doped semiconductor than to the correlated electron materials on which we wish to focus here. Nonetheless some of the scientific excitement has focused on superconductivity (Reyren *et al.*, 2007) (exhibited also when bulk STO is lightly doped by conventional dopants) and Anderson localization metal-insulator transitions (Caviglia *et al.*, 2008). Reports or theoretical suggestions of other

correlation phenomena in this system including charge ordering (Pentcheva and Pickett, 2007) and magnetism (Ariando *et al.*, 2011; Bert *et al.*, 2011; Brinkman *et al.*, 2007; Li *et al.*, 2011; Okamoto *et al.*, 2006) have also appeared; we return to the issue in section IV and in the conclusions.

This Colloquium illustrates the essential features that make TMO-based heterostructures an appealing discovery platform for emergent properties. The guiding principle is that strong electronic correlations in combination with the access to new symmetries and electronic band structures provided by oxide interfaces can activate new electronic properties formerly “hidden” in bulk compounds. We illustrate this principle with a few selected examples, showing how charge redistributes, magnetism and orbital polarization arises and ferroelectric order emerges from heterostructures comprised of oxide components with nominally contradictory behavior. For example, interfaces may be metallic, magnetic, or ferroelectric even though in bulk form the constituent materials are insulating, non-magnetic, or simple dielectrics. We conclude by articulating open challenges and opportunities in the field, in particular, how to translate the new understanding of when emergent phases arise into control of novel behavior by design at oxide interfaces, and the manipulation of these states by suitable mechanical, electrical or optical boundary conditions and excitations.

II. ANATOMY OF AN OXIDE INTERFACE

The formation of a coherent perovskite oxide heterointerface, as shown in Figure 1, provides a remarkable correlated electron “playground.” It brings different transition metal cations with their localized d electron physics and interacting charge, spin and lattice degrees of freedom into intimate contact in a tunable crystalline environment.

The key structural features of transition metal oxides relate to the coordination geometry of the metal ions and the metal-oxygen-metal bond angles. These determine magnetic exchange interactions (Anderson, 1950; Goodenough, 1955; Kanamori, 1965) and electronic bandwidths (Eng *et al.*, 2003), thereby controlling the electronic and magnetic ground states. Structural and electronic changes across an interface can act to stabilize previously unanticipated phases of matter (Okamoto and Millis, 2004).

Consider for example a multilayer heterostructure comprised of alternating blocks of the metallic ferromagnet $\text{La}_{2/3}\text{Ca}_{1/3}\text{MnO}_3$ (LCMO), and the high-temperature cuprate superconductor $\text{YBa}_2\text{Cu}_3\text{O}_{7-x}$ (YBCO) sketched in Figure 1. The interface brings several crucial structural effects. The first is a coordination mismatch. LCMO is a three-dimensional perovskite (AMO_3 stoichiometry) with corner-connected MnO_6 octahedra that may be described by interleaving alternating (La,Ca)O and MnO_2 layers along [001]. In contrast, YBCO is a two-dimensional

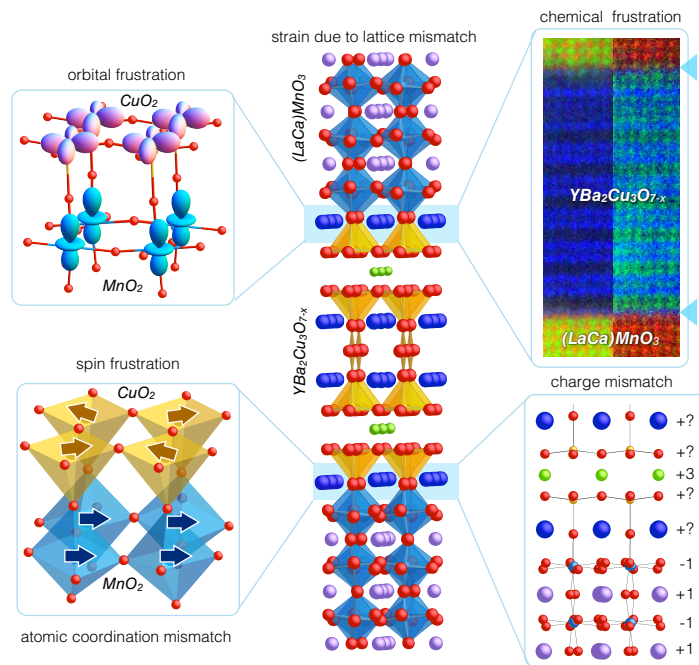


FIG. 1 Anatomy of an oxide heterointerface: an illustration showing the interplay between different degrees of freedom (charge, spin, and lattice) at a coherently grown interface between ferromagnetic $\text{La}_{2/3}\text{Ca}_{1/3}\text{MnO}_3$ and superconducting $\text{YBa}_2\text{Cu}_3\text{O}_{7-x}$. The electron micrograph is reproduced from Ref. [Chien et al., 2013](#), where each color represents a different chemical species.

oxide with four- and five-fold coordinated Cu cations. The layered cuprate structure may be considered as a derivative of perovskite, which partly facilitates coherent growth of the heterostructure. But unlike LCMO, YBCO displays an ordered network of oxygen vacancies accommodated by the valence preferences of Cu: One oxygen atom is removed from every third (001) YO plane to produce the square pyramidal CuO_5 coordination, then on every third CuO_2 layer, vacancies order along [100], producing the square planar CuO_4 coordination. Thus a “coordination mismatch” arising from the change from the 6-fold coordination of the Mn to the lower coordination of the Cu ([Figure 1](#), lower left) occurs at the interface. As a result, a set of CuO chains (*i.e.* charge reservoir) is missing from the interfacial YBCO unit cell to maintain a perovskite-like sequence $\dots\text{MnO}_2\text{--BaO--CuO}_2\dots$ across the junction ([Chien et al., 2013](#); [Zhang et al., 2009](#)).

Coherent epitaxial growth also produces an intrinsic strain mismatch arising from the different equilibrium lattice constants ([Figure 1](#), center). The atomic structure at the heterointerface responds to alleviate the strain mismatch through relaxation of the interatomic distances and internal atomic degrees of freedom (for example, rotations or size deformations to the transition metal oxygen polyhedra) in the constituents along the superlattice repeat direction. These new atomic arrangements directly alter the electronic structure. Away from the interface it is characterized by carriers in the d -manifold with orbital symmetries $d(x^2 - y^2)$ (YBCO) and $d(z^2 - r^2/x^2 - y^2)$ for LCMO ([Figure 1](#), upper left), but near the interface

the $d(z^2 - r^2)$ become occupied in the YBCO and acquire more $d(x^2 - y^2)$ character in the LCMO.

In addition to the structural effects, an electronic mismatch occurs. The ferromagnetism in LCMO relies on the cooperative *parallel* alignment of spins from the narrow correlated electronic bands; singlet Cooper pair formation in YBCO, in contrast, relies on *paired* spins with antiferromagnetic interactions. These antagonistic spin interactions (frustration) have been invoked to explain changes in the interfacial magnetization and superconductivity, *e.g.* giant magnetoresistance, the appearance of uncompensated magnetic moment on Cu in CuO_2 plane, and large modulation of ferromagnetic magnetization profile across the heterojunction ([Chakhalian et al., 2006](#); [Hoppler et al., 2009](#); [Peña et al., 2005](#); [Stahn et al., 2005](#)).

The different valence configurations of the cations in the constituent materials of the heterostructure also induce changes in charge density and chemical bonding. In the system shown in [Figure 1](#) (lower right panel) a charge of $\sim 0.2e$ per Cu ion is transferred from Mn to Cu ions across the interface ([Chakhalian et al., 2007](#)). The charge transfer at other oxide interfaces has also been found to exhibit a peculiar asymmetric electronic “roughness” intertwined with an asymmetric interface stacking sequence or an asymmetric chemical roughness ([Chien et al., 2013](#); [Hoffmann et al., 2005](#); [May et al., 2008](#)). The effects from different stacking sequences and electronic roughness remain to be resolved.

To summarize, the following degrees of freedom are highly tunable at an oxide interface and may be exploited

in uncovering new phases:

- Epitaxial strain mismatch owing to differences in equilibrium lattice parameters
- Atomic coordination frustration and cation site preferences
- Ordered spin and orbital states
- Charge flow across the interface (layer dipole discontinuities)
- Chemical frustration and interlayer mixing

The following examples detail how these considerations are made, and the exciting new phases born from the interplay of the correlated electronic and atomic structure across oxide interfaces.

III. CHARGE AT THE INTERFACE

Understanding and controlling the distribution of charge carriers at the interface between dissimilar semiconductors is one of the pivotal developments of modern microelectronics (Gertner, 2013) important both for devices and as a crucial platform for discovery of remarkable physical phenomena including integer and fractionally quantized Hall effects as well as spin-Hall and other spintronic phenomena.

In conventional semiconductor heterojunctions the basic physics is driven by the difference in work-function, which causes charge transfer across the boundary to equalize chemical potentials. The work-function difference may be manipulated by a process known as δ -doping (Harris *et al.*, 1991; Schubert, 1990), in which a layer of ions is implanted in a plane at some distance from the interface. An additional advantage of δ -doping is that the placement of the dopants at some distance from the interface minimizes the effects of randomness in the dopant positions. δ -doping is now widely used to produce two dimensional electron gases (2DEGs) confined to the proximity to the interface (*e.g.* GaAs/AlGaAs).

The interest in using TMO to explore similar physics was motivated by two observations (Ahn *et al.*, 2003, 2006): (i) in oxides, the accessible carrier density is expected to be orders of magnitude higher than that of semiconductors ($\geq 10^{20} \text{ cm}^{-3}$), and (ii) the Thomas-Fermi screening length is expected to be much shorter, so the charges may be confined to within $<1\text{-}2 \text{ nm}$ of the interface, a factor of $5 - 10$ shorter than the $\sim 10 \text{ nm}$ length characteristic of semiconductor junctions. However, the current intense effort in material synthesis, theory, and device fabrication of oxide interfaces is motivated mainly by the known sensitivity of the correlated electron properties of transition metal oxides to the d -band filling (Armitage *et al.*, 2010; Basov, 2005; Dagotto *et al.*, 2001; Lee and Wen, 2006; Mackenzie, 2003; Ovchinnikov, 2003; Tokura, 2006; Tokura and Tomioka, 1999). The discovery

of an interface-based method of carrier doping has revived the idea of tailoring the materials electronic properties and creating novel quantum states not easily attainable in the bulk counterparts. The basic idea (analogous to that motivating δ -doping) is to explore electronic and magnetic phases without the hindering effects of chemical disorder inherent in the conventional solid state chemistry methods of changing carrier concentration.

During the past several years, extensive experimentation has established that perovskite-based heterostructures are particularly susceptible to interlayer charge redistribution derived from the incompatibilities illustrated in Figure 1 making them ideal candidates to explore such possibilities (Bibes *et al.*, 2011; May *et al.*, 2009; Ohtomo and Hwang, 2004; Ohtomo *et al.*, 2002; Okamoto and Millis, 2004).

A. Interface Doping of a High- T_c Superconductor

To illustrate the inherent interest of charge reconstruction on interfacial states, we discuss as one of many possible examples the recent progress on cuprate/manganite heterointerfaces. Macroscopically it has been established that the introduction of a ferromagnetic (La,Ca)MnO₃ manganite layer into the heterostructure with an optimally doped YBCO cuprate triggers a *suppression* of the superconducting transition temperature accompanied by a reduced ferromagnetic Curie temperature (Driza *et al.*, 2012; Holden *et al.*, 2004; Hoppler *et al.*, 2009; Kalcheim *et al.*, 2011; Peña *et al.*, 2004; Satapathy *et al.*, 2012; Sefrioui *et al.*, 2003). In a recent set of experiments (Figure 2), L -edge polarized resonant X-ray absorption spectra taken at the Mn and Cu edges reveal the presence of a chemical shift implying a flow of electronic charge across the interface of about $\sim 0.2e$ per Cu atom (Chakhalian *et al.*, 2007; Chien *et al.*, 2013). The depleted electrons from MnO₂ layer are directly transferred to the CuO₂ planes, unbalancing the charge distribution between the atomic CuO₂ layers and the CuO chain charge reservoir block. The average Mn valence also increases from the as-grown value ($\text{Mn}^{+3.33}$) to around 3.5, indicative of covalent bond formation across the Mn–O–Cu interface.

The charge transfer across the interface from the Mn to Cu ions induces a major reconstruction of the d -orbital occupancies and frontier orbital symmetries in the interfacial CuO₂ layers (Chakhalian *et al.*, 2007, 2006). In particular, the Cu $d_{3z^2-r^2}$ orbital, which is fully occupied and electronically inactive in the bulk cuprates becomes active at the interface (Figure 2b). At the same time charge transfer is observed in the presence of enhanced covalent chemical bonding across the interface, the Cu cations from the nominally antiferromagnetic CuO₂ plane acquire an uncompensated magnetic moment (Figure 2c), attributed to spin canting of the local moments on the interfacial Cu cations.

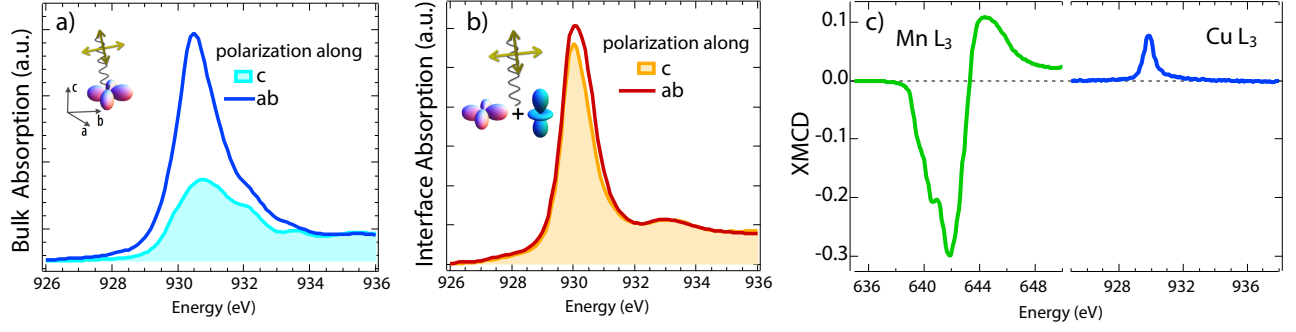


FIG. 2 Electronic structure of Cu in a $\text{La}_{2/3}\text{Ca}_{1/3}\text{MnO}_3/\text{YBa}_2\text{Cu}_3\text{O}_{7-x}$ heterostructure, determined from X-ray Linear Dichroism (XLD) and X-ray Magnetic Circular Dichroism (XMCD) measurements. Panels (a,b): XLD spectra taken on the Cu L_3 -edge at temperature $T = 15$ K with the electric-field vector $\mathbf{E} \parallel ab$ plane and $\mathbf{E} \parallel c$ plane, taken in bulk (panel a) and interface (panel b) sensitive modes. The main peak (“white line”) in (b) is shifted towards higher energies, indicating a lower charge state of Cu at the interface. Panel (c): XMCD spectra measured at the Cu and Mn L_3 edges in (c) recorded at $T = 15$ K in a 5 T applied magnetic field demonstrating that the interfacial copper cations exhibit a non-zero ferromagnetic local moment, whereas in bulk the antiferromagnetic coupling leads to a net magnetization of zero.

Initial studies of the interplay between the ferromagnetic and superconducting order parameters used synchrotron based X-ray and neutron reflectivity experiments. However these tools were unable to clearly resolve the length scale of interactions at the boundary between the two phases. Very recently, the issue has been

addressed by use of cross-sectional scanning tunneling microscopy (XSTM) together with atomic-resolution electron microscopy (EELS/STEM). These methods enable direct observation of the charge distribution and the corresponding spatial scale for the buried interface (Chien *et al.*, 2013). Figure 3 shows the spatially resolved dI/dV spectra, which provided the first direct evidence that the length scale for charge transfer between YBCO and LCMO has an upper limit of <1 nm, and that the spatial broadening of the electronic transition is commensurate with the rougher interface. This result sets a fundamental upper limit on the charge-transfer length scale in the YBCO/LCMO system, ruling out a class of theories based on long-range proximity effects (Hoffmann *et al.*, 2005). In addition to the established X-ray and neutron based probes, this powerful characterization technique provides a useful tool to achieve a microscopic direct space understanding of the electronic structure across correlated oxide interfaces.

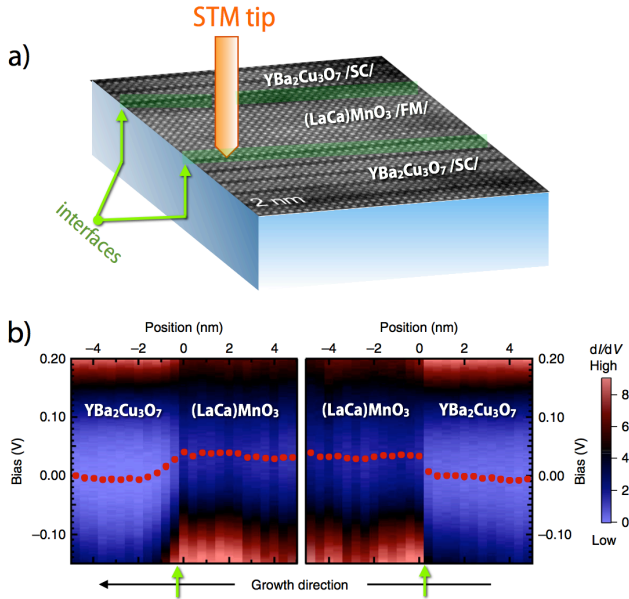


FIG. 3 Panel (a): Schematic of cross-sectional scanning tunneling microscopy (XSTM) measurements performed on an LCMO/YBCO superlattice grown on a Nb-doped STO substrate. Panel (b): Data reproduced from Ref. Chien *et al.*, 2013. The spatial evolution of the dI/dV spectra averaged across the two identically terminated heterointerfaces reveals that the electronic transition is more abrupt for the bottom interface (right arrow) than the top, broader, interface (left arrow). The red dots represent the voltage of the minimum in the density of states.

B. Additional considerations

The complex behavior occurring at the LCMO/YBCO interface highlights the need to develop a clear language and set of concepts to describe interface electronic physics in correlated oxides. The inherently many-body nature of the correlated interface raises fundamental questions, in particular of the applicability of the ideas, formulae, and language devised for semiconductor interfaces where a single-particle description works well. Pioneering work of Oka and Nagaosa (Oka and Nagaosa, 2005) showed via density matrix renormalization group calculations of a one dimensional model system (in essence the one dimensional Hubbard model with a spatially varying interaction parameter and band bottom) that the standard concepts of band bending and interface dipole apply, albeit with

some modifications, as long as the conduction and valence bands are replaced by lower and upper Hubbard bands.

A growing body of literature builds on this work, using the concepts of band bending, Schottky barriers, and depletion layer creation borrowed from semiconductor physics (Hikita *et al.*, 2009; Yajima *et al.*, 2011), as well as more involved approaches, which unite Poisson-Schrödinger electrostatics with Mott-Hubbard physics (Charlebois *et al.*, 2013; Lee and Macdonald, 2006; Okamoto and Millis, 2004). Correlation physics is shown to lead to *quantitative* changes in the spatial confinement of carriers near interfaces (Lee and Macdonald, 2006, 2007; Okamoto and Millis, 2004), including the possible formation of extended depletion regions of zero compressibility (so-called ‘Mott plateaus’)(Charlebois *et al.*, 2013; Lee and Macdonald, 2006). Other theoretically proposed possibilities *unattainable with semiconductor junctions*, include a spontaneously emerging quantum-well structure when an electron-doped Mott-Hubbard insulator is coupled to a normal metal with a large work-function. Following the same line of reasoning, in a *p-n* junction between two correlated insulators the local Mott gap collapses giving rise to a 2DEG (Charlebois *et al.*, 2013).

With few exceptions (Jin *et al.*, 2011), current experimental attention has focussed on interfaces such as that between the two band insulators LaAlO_3 and SrTiO_3 . In most of these situations the carriers are introduced via the polar catastrophe mechanism (Mannhart *et al.*, 2008); the maximum sheet carrier density is 0.5 per in-plane unit cell and this carrier density is typically distributed (Mannhart *et al.*, 2008; Okamoto *et al.*, 2006) over several unit cells away from the interface, leading in general to volume carrier densities far below the Mott value of one per unit cell. Density functional plus Hubbard U calculations (Pentcheva and Pickett, 2007) indicate that a charge ordered phase in which the entire polar catastrophe charge density is in the first interface layer may be possible, but these suggestions have not yet been confirmed by experiment or beyond-DFT methods. One very interesting potential exception is the work of Moetakef *et al.*, 2012 on $\text{GdTiO}_3/\text{SrTiO}_3$ heterostructures, where a nontrivial insulating phase was observed when two layers of SrTiO_3 were sandwiched between thick sheets of GdTiO_3 . It has been explained by Chen, Lee and Balents in terms of a novel ‘Mott dimer’ phase (Chen *et al.*, 2013), where the carrier density is far below the one electron per transition metal ion value needed for Mott physics; nonetheless many theoretical predictions suggest alternative avenues for emergent properties to arise and warrant experimental investigation.

Additional issues beyond conceptual approaches to interface control arise. The length scales in correlated oxides are typically very short, so the details of the interface may be more important than in conventional semiconductors. A local picture is needed, which is able to address the formation of chemical bonds across the junction, differing elec-

tronegativities of transition metal ions, changes in both crystal field energies and Madelung potentials, and polarity effects (Biscaras *et al.*, 2012; Garcia-Barriocanal *et al.*, 2013; Herranz *et al.*, 2007; Hotta *et al.*, 2007; Ohtomo and Hwang, 2004; Park *et al.*, 2013b; Salluzzo *et al.*, 2013; Savoia *et al.*, 2009; Sing *et al.*, 2009; Takizawa *et al.*, 2009; Zhong *et al.*, 2010).

A further complication is that while many correlated oxides are reasonably well described by the Mott-Hubbard picture on which the above-cited works are based, some important functional TMO are *charge-transfer* compounds (Imada *et al.*, 1998; Khomskii and Sawatzky, 1997; Zaanen *et al.*, 1985). The role of the lower Hubbard band in these materials is usurped by the ligand states (typically oxygen $2p$), thus implying a very different physical character for the doped holes (mainly in oxygen levels) and doped electrons (mainly in transition metal d -levels). As a result, the alignment of the oxygen levels across the interface becomes crucial.

For all of the materials discussed in this paper, theoretical treatments which go beyond the simple Hubbard model, including chemically realistic structures and energetics on the same footing as correlation effects, are needed, as are experimental investigations of systems with higher electron densities and complete control over cation and oxygen stoichiometry.

IV. CONTROL OF MAGNETISM WITH OXIDE HETEROSTRUCTURES

Long range magnetic order in transition metal oxides usually arises from a combination of local moment formation on the transition metal site and inter-site coupling via the oxygen sublattice. Heterostructures offer an opportunity to generate new magnetic states by manipulating both the moment formation and the nature of the inter-site coupling. As examples, we note that the paramagnet LaCoO_3 can be converted to a ferromagnetic (FM) material by tensile epitaxial strain, which changes the material from a low-spin to a high-spin state (Freeland *et al.*, 2008; Fuchs *et al.*, 2007; Park *et al.*, 2009; Rondinelli and Spaldin, 2009). On the other hand, bulk antiferromagnetic (AFM) EuTiO_3 can be converted to a ferromagnetic insulator under modest tensile strains (Lee *et al.*, 2010). Another notable example is the comprehensive study by Seo *et al.*, 2010, which examined three-component $\text{SrRuO}_3/\text{manganite}/\text{SrRuO}_3$ heterostructures. These authors found strong compressive strain causes relative FM alignment of magnetization in the heterostructure layers, while tensile or weak compressive strain favors AFM alignment of neighboring layers.

This sort of control over local magnetization in thin film geometries is of potential utility for oxide electronics and spintronic applications, including magnetic memory and sensing (Bibes *et al.*, 2011). For example, electromechani-

cal coupling via a piezoelectric material can be used to control the orientation and strength of the magnetization by tuning the lattice parameters of the heterostructure through an applied electric field (Dekker *et al.*, 2011). Here, we focus on going beyond strain control to make use of the broken symmetry at the interface between two dissimilar materials to generate unique spatially structured magnetic states.

A. Creating Novel Magnetic States at Interfaces

One approach to manipulating magnetism involves interfacial charge transfer in heterostructures created from an antiferromagnetic *insulator* and a paramagnetic *metal* (Freeland *et al.*, 2010; Takahashi *et al.*, 2001; Yordanov *et al.*, 2011). The choice of materials in this case was determined by two key factors: first, creating moments from a material without any propensity to moment formation, *i.e.*, zero moments, is difficult. It is therefore reasonable to begin then by choosing a system with a large local moment such as CaMnO_3 with $3\mu_B/\text{Mn}$, which in bulk is a G-type (conventional two-sublattice Néel) antiferromagnet. However, modest electron doping of this material leads to strong ferromagnetic (FM) correlations (Neumeier and Cohn, 2000). In a quantum-well heterostructure in which a paramagnetic metal (in this case CaRuO_3) is confined between two thick layers of CaMnO_3 one may expect that charge transfer from the metal to insulating CaMnO_3 will lead to interfacial doping and thus ferromagnetism.

Theoretical studies substantiate this argument and find that a charge of approximately $0.1e$ per interface unit cell leaks across the interface and is confined within ~ 1 unit cell at the $\text{CaRuO}_3/\text{CaMnO}_3$ interface (Nanda *et al.*, 2007). Although the magnitude of the charge leakage is small, it has a significant impact on the antiferromagnetic order in the CaMnO_3 , providing a mechanism for spin canting which yields large ferromagnetic moments at the interface (Freeland *et al.*, 2010; Takahashi *et al.*, 2001; Yordanov *et al.*, 2011). To validate this concept a study of the spatial distribution of the magnetism was carried out using X-ray resonant magnetic scattering (XRMS) at the Mn *L*-edge (Freeland *et al.*, 2005; Kavich *et al.*, 2007). Figure 4 shows the large XRMS signal, and that it deviates from anticipated bulk G-type AFM state, which shows no ferromagnetic component to the magnetic moment under identical strain conditions indicating that the ferromagnetism emerges from the interface (Freeland *et al.*, 2010). By fitting this signal as a function of incident angle, the extent of the magnetic polarization away from the interface was found to extend over several unit cells in contrast to the length of one unit cell predicted by theory (Nanda *et al.*, 2007). The observed longer length scale of the magnetization profile discrepancy may be due to magnetic polarons, which are known to exist in lightly-doped CaMnO_3 (Chiorescu *et al.*, 2007), but such interfacial

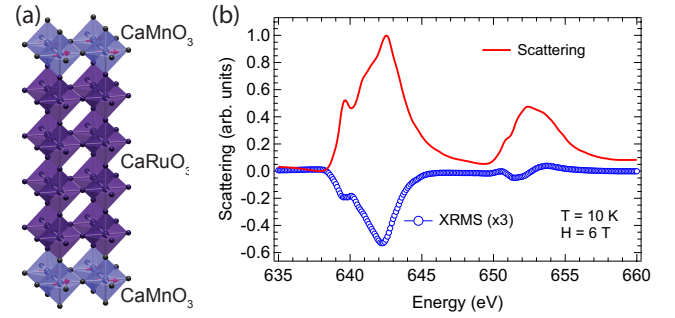


FIG. 4 Panel (a): Schematic crystal structure showing canted spins (pink arrows) within the MnO_6 octahedra of CaMnO_3 at the interface of the quantum-well structure with metallic CaRuO_3 . The canting arises from electron transfer owing to the ohmic contact. Panel (b): X-ray resonant magnetic scattering data showing a large magnetic signature arising from the FM alignment of spins at the interface in the presence of a magnetic field (see associated data in Ref. Freeland *et al.*, 2010).

polarons have not explicitly investigated theoretically.

B. Other Routes to Interface Magnetism

Strain and layer sequencing can offer additional handles to manipulate the interfacial magnetic state in the $\text{CaRuO}_3/\text{CaMnO}_3$ system (He *et al.*, 2012). The link between the metallic layer and magnetism is best illustrated by studying superlattices where the metallic layer undergoes a metal-to-insulator transition when the dimensionality is reduced in the ultra-thin layer, and correspondingly the magnetism disappears (Grutter *et al.*, 2013). One can use this understanding and exploit it to design new functional materials and there are many possibilities that exist within the perovskite family which can be combined to seek new types of magnetic states (Bhattacharya *et al.*, 2008; Gibert *et al.*, 2012; Hoffman *et al.*, 2013; Smadici *et al.*, 2007). For example, many antiferromagnets have ordering temperatures well-above room temperature, so one could extend this concept to create interfacial insulating *ferrimagnets* that operate at high temperature (Ueda *et al.*, 1998). Another possibility is to make use of the spatially localized magnetic state in proximity to a metallic layer to create a spin-polarized 2DEG (Nanda and Satpathy, 2008). More broadly, one could create heterostructures with two magnetic materials, and use the competition towards different collectively ordered magnetic states in addition to structural incompatibilities to generate a plethora of interesting and potentially spatially varying magnetic phases. These are but a few of the magnetic possibilities which remain to be uncovered at oxide heterointerfaces, chosen to highlight the large phase space still available for exploration and the opportunities available to connect with materials theory in the rational search for new magnetic systems.

V. INTERFACIAL CONTROL OF ORBITAL POLARIZATION

A. The Case of Rare-earth Nickelates

The orbital configuration, *i.e.*, the distribution of the d -electrons over the available crystal field levels, plays an important role in the formation of strongly correlated ground states in transition metal oxides (Tokura and Nagaosa, 2000). In general, orbital configurations are closely linked to structure and may therefore be manipulated at interfaces. Here we discuss these issues specifically for the orthonickelate perovskites $R\text{NiO}_3$, where R is a trivalent cation from the lanthanide series, but the ideas can be extended to other AMO_3 systems.

The original and decade later renewed interest in nickelates arose from the possibility of generating a cuprate-like electronic and orbital configuration in a copper-free system (Anisimov *et al.*, 1999; Chaloupka and Khaliullin, 2008; Hamada, 1993; Lee and Pickett, 2004; Poltavets *et al.*, 2010). The basic idea is that in bulk $R\text{NiO}_3$ the Ni is octahedrally coordinated, with only small deviations from cubic (O_h) symmetry. Further, formal valence considerations indicate that the nominally Ni^{3+} cation is in the low-spin d^7 configuration, with the t_{2g} states ($d_{xy,xz,yz}$) filled and one electron in the two-fold degenerate e_g -symmetry ($d_{3z^2-r^2,x^2-y^2}$) Ni d -levels. Low-spin d^7 is a first-order Jahn-Teller configuration, with a susceptibility to bond distortions which break the cubic point symmetry and are enhanced by correlation effects. It was thus expected that modest perturbations would split the e_g levels, leaving an effective one-band configuration where the electron is fully confined to a single orbital.

The degree to which an electron occupies two different m_{l1} and m_{l2} orbitals can be quantified as an *orbital polarization*

$$P_{l_1 m_{l1}, l_2 m_{l2}} = \frac{n_{l_1 m_{l1}} - n_{l_2 m_{l2}}}{n_{l_1 m_{l1}} + n_{l_2 m_{l2}}},$$

where $n_{l_1 m_{l1}}$ and $n_{l_2 m_{l2}}$ are the occupancies of the $|l_1 m_{l1}\rangle$ and $|l_2 m_{l2}\rangle$ states (Han *et al.*, 2010), with orbital quantum number l_i and magnetic quantum number m_{li} , respectively. For the rare-earth nickelates, the relevant orbital polarization arises from the $n_{x^2-y^2}$ and $n_{3z^2-r^2}$ occupancies, and a fully polarized state $P = 1$ would be indicative of a single band electronic structure.

Something akin to this effect occurs in many members of the “colossal” magnetoresistance manganites, where the basic configuration is a high-spin d^4 configuration and similarly a Jahn-Teller ion that can be manipulated with strain (Tokura and Nagaosa, 2000). Hubbard-model calculations further indicated that the single-band physics was very likely to appear (Hansmann *et al.*, 2009); however, more realistic *ab-initio* calculations indicate that the actual electronic configuration for Ni is in the high-spin d^8 state with a hole on the oxygen ($d^8\bar{L}$) (Han *et al.*, 2010).

Since the high-spin d^8 configuration has one electron in each of the two e_g orbitals, it is significantly less susceptible to undergoing Jahn-Teller distortions, suggesting that it would be more difficult than initially expected to achieve the desired degree of orbital polarization, even in the correlated case (Han *et al.*, 2011). Studies of the dependence of orbital polarization on the different flavors of structural symmetry-breaking (Cammarata and Rondinelli, 2013) is thus of great experimental interest and is a stringent test of the theory.

B. Manipulating Orbitals in $R\text{NiO}_3$ Heterostructures

Advances in high-quality growth of nickelates over the past few years mean that we are now in a position to test these predictions (Boris *et al.*, 2011; Bruno *et al.*, 2013; Eguchi *et al.*, 2009; Hwang *et al.*, 2013; Liu *et al.*, 2010b; May *et al.*, 2010; Scherwitzl *et al.*, 2010; Tsubouchi *et al.*, 2008). The basic experimental approach is to use a combination of quantum confinement, achieved by fabricating ultra-thin layers of TMO sandwiched between layers of wide-gap insulators, and epitaxial strain, obtained by varying the substrate material, to break the octahedral symmetry. Advanced x-ray techniques are then used to estimate the resulting changes in orbital occupancies.

However, *ab-initio* calculations based on density functional theory indicate that the contribution of strain to octahedral symmetry breaking is not completely intuitive (see Rondinelli *et al.*, 2012 and references therein). In particular, a considerable degree of compression or tension can be accommodated by octahedral rotations, without necessarily changing the local point symmetry significantly since the NiO_6 units are highly flexible (Chakhalian *et al.*, 2011). Furthermore, quantum confinement may be affected by the chemistry of the insulating layer, with different degrees of polarization found for different choices of wide-gap insulator (Han *et al.*, 2010).

At present, the experimental results are not completely consistent with each other or with theory. For example, examination of the Ni L_2 edge indicated an $\sim 5\%$ orbital polarization for a single unit-cell of LaNiO_3 subject to tensile strain (Freeland *et al.*, 2011) and no orbital polarization for compressive strain. Other measurements employing an orbital reflectometry technique on four unit cell films also observed a similar non-zero interfacial polarization for tensile strain (Benckiser *et al.*, 2011; Frano *et al.*, 2013). Recent studies have indicated it is possible to increase the orbital polarization up to 25% through judicious optimization of high tensile strain states and alternative spacer materials (Wu *et al.*, 2013); the latter had been shown theoretically to play a considerable role in obtaining the targeted orbital polarization levels (Han *et al.*, 2010).

C. Open Questions in Orbital Control at Interfaces

All experiments agree though that the degree of orbital polarization observed in actual superlattices is small compared to that needed to achieve a fully orbital polarized Ni e_g^1 state. The main challenge is to then build the framework to understand how to create fully orbital polarized states in oxide heterostructures.

One important facet of this problem has to do with strain and symmetry. For example, LaNiO_3 has rhombohedral symmetry in the bulk which actually disfavors a uniaxial Jahn-Teller distortion (Carpenter and Howard, 2009). NdNiO_3 , on the other hand, is orthorhombic which allows such a distortion without large energetic penalties. Recent studies by Tung *et al.*, 2013 show that the nickelate films maintain to some extent the symmetry of the bulk, which, due to the connection between compatible lattice distortions and crystal symmetry, directly influences the ability to orbitally polarize the $3d$ -states even under large strains.

With this understanding, one may be able to choose the proper bulk symmetry of the TMO to be used in the heterostructure to build in larger orbital polarizations in NdNiO_3 by coupling strain with the interfacial covalency effect discussed above and interfacial proximity effects (Aso *et al.*, 2014; Moon *et al.*, 2014). Even for the case of NdNiO_3 films, however, the orbital polarization is still insufficient to create a fully polarized state (Tung *et al.*, 2013). This is largely due to the energy scale mismatch between elastic strain (~ 100 meV) and the bandwidth (on order of several eV), and the overall tendency to orbital polarization is further reduced by the $d^8\bar{L}$ character of the Ni^{3+} state.

Small orbital polarizations have also been observed even in the case of the Jahn-Teller active manganites (Aruta *et al.*, 2006; Pesquera *et al.*, 2012; Tebano *et al.*, 2008), which indicates that this balancing of drastically different energy scales is difficult even in systems that prefer orbital order. A potential solution is to create interfaces with large symmetry mismatch due to lattice topology or by combination of dissimilar crystal field environments.

Consider for example bulk oxides with large orbital polarization such as the cuprates (Chen *et al.*, 1992; Nücker *et al.*, 1995) and Ruddlesden-Popper (layered-structure) nickelates (Kuiper *et al.*, 1998; Pellegrin *et al.*, 1996) as a starting point. In these materials, the large orbital polarization arises from the strongly asymmetric crystal (ligand) field of the layered structure. As was discussed above for the LCMO/YBCO heterointerface, oxide interfaces can be harnessed to ‘undo’ orbital polarization, but there is no reason why the converse should not also be possible. This offers a real opportunity in the area of matching systems with drastically different symmetries to create orbital states at the interface.

Orbital control can also be used to modulate strongly correlated states. Strain very effectively controls the

metal-insulator transition (MIT) for NdNiO_3 thin films (Liu *et al.*, 2010a, 2013), but the underlying mechanism is not fully understood. Using quantum confinement when the layer dimensions approach the atomic limit, it was observed that orbital polarization under compressive strain tends to favor a metallic state while quantum confinement caused a re-emergence of a MIT through the interfacial reduction of the orbital polarization (Liu *et al.*, 2012a). A similar connection was recently observed in the case of VO_2 thin films (Aetukuri *et al.*, 2013), where the decrease in the MIT temperature was correlated with strain driven polarization of the V t_{2g} orbitals. The potential use of strain in combination with symmetry mismatch to tune between correlated metallic and insulating phases is an important issue warranting further investigation.

VI. FERROELECTRIC HETEROSTRUCTURES FROM NONFERROELECTRIC BULK OXIDES

The electrically switchable polarization of ferroelectrics (FE) allows their integration in random access memories (FE-RAM), electro-optical devices, sensing microsystems, active vibration control and surface acoustic wave systems, to high frequency devices (Setter *et al.*, 2006). The main challenges for future FE-RAM scaling, however, is that the FE dielectric thickness must be reduced to fit within the required device area while maintaining sufficient reproducibility and signal margins for sense amplifier differentiation between a ‘0’ and ‘1’ data state (Wu *et al.*, 2010). Furthermore, non-destructive *magnetic* sensing of *electric* polarization, enhanced miniaturization and increased packaging density in magnetoelectric materials (ME) (Eerenstein *et al.*, 2006; Fiebig, 2005; Ramesh and Spaldin, 2007; Velev *et al.*, 2011) would enable the realization of four-state logic in a single device (Bibes and Barthelemy, 2008; Khomskii, 2009).

The conventional approach for realizing strong ME materials, i.e., where there is strong coupling between the primary electric and magnetic polarizations, uses naturally occurring materials possessing primary ferroic orders, namely ferroelectricity and ferromagnetism. Such materials not only are rare, but often suffer from weak coupling between the spin and charge degrees of freedom (Eerenstein *et al.*, 2006).

Recent advances in atomic layer epitaxy now enable the design and fabrication of heterostructures with atomically flat interfaces that can support new forms of ferroelectricity (Bousquet *et al.*, 2008; Mulder *et al.*, 2013; Rondinelli and Fennie, 2012) and magnetoelectric coupling owing to interfacial interactions among electronic spins, charges, and orbitals (Wu *et al.*, 2010). A promising avenue to pursue in the search for new materials with emergent ferroelectricity and a strong magnetic field dependence of the electric polarization exploits an superlattice structure with broken inversion symmetry, which results from

being constructed from three distinct layers (Lee *et al.*, 2005; Warusawithana *et al.*, 2003). The ‘tri-color’ layering lifts inversion symmetry – a prerequisite for an electric polarization – whereas epitaxial strain applied to the heterostructure can promote the formation of electrically and magnetically tunable polarizations, even in the absence of ferroic components (Hatt and Spaldin, 2007; Tokura, 2007).

Using a combination of complementary experimental probes, magnetoelectricity was recently demonstrated in artificial tri-layer heterostructures consisting solely of dielectric antiferromagnetic oxides (Figure 5a). Laser molecular-beam epitaxy was used to create the heterostructure comprising alternating LaMnO_3 , SrMnO_3 , NdMnO_3 layers on a SrTiO_3 substrate. Rogdakis *et al.*, 2012 report the emergence of ferroelectricity below 40K (Figure 5c) and it was found to depend on the number of NdMnO_3 layers n in the superlattice (Figure 5d). Interestingly, the authors observed slim loop-like polarization–electric (P - E) field hysteresis, with an extended tail of the polarization above the ferroelectric transition temperature and a thermal hysteresis between zero-field-cooled and field-cooled measurements. Such features are typical of relaxor ferroelectrics and were attributed to interface effects (Rogdakis *et al.*, 2012). We note that this dielectric relaxation also leads to differences in the magnitudes of the measured polarization obtained from the P - E loop and the pyrocurrent measurement, which might also be affected from the challenges in characterizing the dielectric properties of ultrathin film oxides with techniques commonly used for bulk single crystals. Nonetheless, the magnetoelectric coupling resulted in 150% magnetic modulation of the electric polarization, demonstrating how heterostructuring multiple compounds together to lift inversion symmetry in superlattices is an avenue to create new functionalities.

First-principles density functional calculations indicated that broken space inversion symmetry and mixed valency, arising from the heterostructure geometry (cation layer sequence) and interfacial polar discontinuity, respectively, is responsible for the observed behavior. In particular, the formal charge layering of the LaMnO_3 and NdMnO_3 components at the interfaces with SrMnO_3 give rise to a charge discontinuity, leading to electron transfer and cooperative off-centering of the cations. The A cation layering leads to a pattern of Mn and A-cation displacements along the superlattice normal growth direction that lift inversion symmetry and therefore produce the macroscopic electric polarization. We note that the ferroelectric relaxor behavior could not be seen from the theoretical results, which capture the static and cation ordered zero-temperature behavior.

This work demonstrates yet another fascinating example of emergent functionality exhibited in heterostructures. The ability to lift inversion symmetry and independently tune spin order allows the design of many more materials

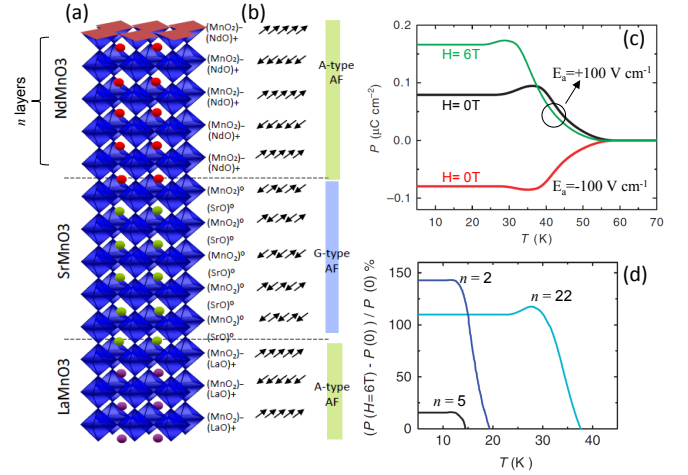


FIG. 5 Ferroelectric and magnetoelectric properties of $[(\text{NdMnO}_3)_n/(\text{SrMnO}_3)_n/(\text{LaMnO}_3)_n]_m$ superlattice, where (n,m) denotes the specific superlattice structure (a) Schematic $[(\text{NdMnO}_3)_5/(\text{SrMnO}_3)_5/(\text{LaMnO}_3)_5]_8$ superlattice on single-crystalline SrTiO_3 substrate with the metal–oxygen octahedra and A cations emphasized. The arrays of the arrows in (b) represent the corresponding antiferromagnetic spin arrangements for each component of the heterostructure. (c) Temperature (T) dependence of the electric polarization (P) measured in a superlattice of period (22,2) using the pyroelectric technique for a typical electric field (E_a) of $+100 \text{ V cm}^{-1}$ (black curve) and -100 V cm^{-1} (red curve) applied perpendicular to the plane of the superlattice layering. The temperature-dependent electric polarization under a magnetic field $H=6 \text{ T}$ applied parallel to the plane of the superlattice layering (green curve) reveals strong magnetoelectric coupling. (d) Normalized relative change in the electric polarization at fixed electric and magnetic fields for various superlattices. Figure adapted from Ref. Rogdakis *et al.*, 2012.

with multifunctional behavior (Gou and Rondinelli, 2014; Puggioni and Rondinelli, 2014). One may exploit these systems to engineer devices from artificial low-dimensional materials exhibiting novel tunable functions distinct from that of bulk systems.

VII. CONCLUSION

The physics of interfaces between materials exhibiting correlated electronic behaviors including superconductivity, magnetism and ferroelectricity is a rapidly advancing field, situated at the intersection of materials science, solid state chemistry and condensed matter physics. Understanding and exploiting these remarkable systems places extraordinary demands on synthesis, measurement and theory, and the challenge is stimulating remarkable work in all areas. By way of conclusion we highlight challenges and prospects in correlated oxide interfaces.

A. Chemical and structural order

Characterization and control of chemical and structural order is a crucial issue. While research to date has revealed remarkable phenomena, clearly related to properties of theoretically ideal interfaces, effects of disorder are not negligible. The brutally short length scales (often only one or two unit cells) pose strong constraints on materials quality. For example, metal to insulator transitions generically occur in oxide heterostructures when the thickness of the metallic layer becomes of the order of 1-2 unit cells. Systematic dependence on strain (Son *et al.*, 2010), and systematic evolution of electronic structure with thickness (Yoshimatsu *et al.*, 2011) suggest an important intrinsic component, but disorder effects and changes in growth processes on these length scales cannot yet be ruled out as mechanisms. Antisite defects mean that real interfaces are not as sharp as depicted in the idealized sketches shown in this paper, and these defects are not necessarily easy to identify in transmission electron microscopy experiments, which average over columns of order 10^3 atoms. Further, oxygen defects and interstitials play a crucial role in transition metal oxides and oxygen partial pressure during growth and in post-growth annealing of heterostructures clearly affects properties in many cases (Ariando *et al.*, 2011; Nakagawa *et al.*, 2006). Methods to further define and control the actual structure of interfaces are urgently needed. One area of future study, is to couple the insight from *in-situ* studies of oxide film synthesis to that of multiscale theory in order to build a mechanistic understanding of the process by which interfaces are created.

B. Theory

The importance and interest of oxide interfaces for the general issue of the theory of correlated electron materials cannot be overemphasized. Understanding the phenomena at interfaces requires a combination of sophisticated many-body physics (to understand the correlated electron states) and *ab-initio* insights (to understand the implications of the changes in octahedral rotations, atomic coordination, and lattice relaxations). The present state of the theoretical art is a combination of analysis of model systems (in particular the Hubbard model), which cannot easily encode many real materials aspects, in particular the transition-metal/ligand covalence as well as the energetics associated with lattice relaxations and *ab-initio* techniques (especially the DFT+*U* method) which have provided crucial insights but are based on a greatly oversimplified Hartree approximation to the many-body physics and may overemphasize order (Chan *et al.*, 2009; Wang *et al.*, 2012). In particular the status of the DFT+*U* predictions of magnetism (Okamoto *et al.*, 2006) and charge order (Pentcheva and Pickett,

2007) at the LAO/STO interface remains unclear.

The combination of density functional band theory and dynamical mean field theory (DFT+DMFT) is a promising alternative (Kotliar *et al.*, 2006), combining *ab-initio* and many-body physics in a systematic way. However, working implementations of total energy calculations are only now beginning to appear (Park *et al.*, 2013a) and forces cannot yet be computed so structural optimization remains a challenge. More fundamentally, existing implementations for systems in which more than one *d*-orbital is important are based on the single-site approximation, which is believed to become poor in the two dimensional situation relevant to heterostructures.

C. Topological states of matter

Topological insulators (TIs) are a fascinating class of materials in which strong spin-orbit interaction promotes gapless electronic states on the surface (*i.e.* edge states) with the bulk of a material remaining gapped (Fu *et al.*, 2007; Hasan and Kane, 2010; Hsieh *et al.*, 2008; Moore, 2010; Qi and Zhang, 2010). Most of the current TI materials belong to the Bi_2X_3 ($X=\text{Se, Te}$) family. Recently, a new approach has been proposed that is based on superlattices of two (or three) unit cells of a strongly correlated electron perovskite ABO_3 grown *along the [111] direction* combined with a band insulator spacer layer; the resulting heterostructure structurally forms a buckled honeycomb lattice topologically equivalent to that of graphene lattice for the case of three unit cell strongly correlated oxide. Depending on the strength of electron-electron correlations, magnitude of Hund's coupling and inter-site hopping, the proposed heterostructures display potentially rich physics associated with exotic electronic and topological phases (Okamoto, 2013; Rüegg and Fiete, 2011; Rüegg *et al.*, 2012, 2013; Xiao *et al.*, 2011; Yang *et al.*, 2011). At present, the main challenge in experimental realization is the film growth along the [111] direction since for the commonly used substrates, *e.g.* SrTiO_3 LaAlO_3 , NdGaO_3 , YAlO_3 , etc., the (111) structure consists of alternating $\pm 4e$ or $\pm 3e$ charged planes along this direction. The large polar discontinuity generally results in complex surface/interface and electronic reconstructions (Enterkin *et al.*, 2010; Marks *et al.*, 2009), which can act to compensate for the polar mismatch. To date there is limited understanding of thin film nucleation, growth and charge compensation in perovskites along highly polar directions. Very recently the synthesis work in this direction has been initiated (Middey *et al.*, 2012).

D. Oxygen Defect control

While many of the examples discussed above involve oxygen stoichiometric perovskites, the ease of re-

moval/addition of oxygen can also offer opportunities for materials that can be programmed by their chemical environment (Kalinin *et al.*, 2012; Kalinin and Spaldin, 2013). While the role of oxygen vacancies has been explored deeply in the context of catalysis and fuel cells (Adler, 2004), recent work has highlighted the controlled stabilization of related oxygen deficient phases using oxide heterostructures. This is interesting for epitaxial thin film phases such as $\text{SrCoO}_{3-\delta}$ (Jeen *et al.*, 2013a,b) or $\text{La}_{1-x}\text{Sr}_x\text{FeO}_{3-\delta}$ (Xie *et al.*, 2013), which can be reversibly converted between oxygen deficient and stoichiometric phases at low temperatures. Since these phases have drastically different ground states, it offers an interesting path for control of strongly correlated electrons via dynamic anion compositional control. By combining low conversion energy with electrochemical gating of vacancies, such as that seen recently for VO_2 (Jeong *et al.*, 2013) and RNiO_3 (Shi *et al.*, 2013), this approach allows direct control of metal vs. insulating phase as well as possible elements of brain-like (neuromorphic) electronic circuits.

E. Moving beyond the static realm

Up to now, all the properties that have been discussed were limited to the quasi-equilibrium properties, but in the future one should also investigate the *dynamical* degree of freedom to explore the emergence of unique transient states. While the dynamic response for bulk materials has been extensively investigated (Averitt and Taylor, 2002; Basov, 2005), oxide heterostructures offer new possibilities. Recent pump-probe studies of oxide films illustrate the potential for ultrafast strain modulation (Daranciang *et al.*, 2012; Wen *et al.*, 2013), which allows one to manipulate the lattice in a new direction since the film motion is clamped in-plane by epitaxy and can only alter the lattice out of plane. Using this epitaxial constraint allows one to drive the crystalline lattice (symmetry, rotations, etc...) into distinctly different areas of phase space. For example, experiments in manganite thin films showed the emergence of a *hidden* phase that existed only in the dynamic realm (Ichikawa *et al.*, 2011). Moving into the mid-IR region enables direct pumping of lattice modes that can trigger phase transitions (Rini *et al.*, 2007) and was recently used to trigger a metal-insulator transition through dynamic strain created by direct pumping of substrate phonons (Caviglia *et al.*, 2012). Low energy photons in the THz regime can also serve as a dynamic way to drive transitions with ultra-fast electric fields (Liu *et al.*, 2012b). Such experiments have only begun to explore the complex landscape available in the dynamic realm.

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